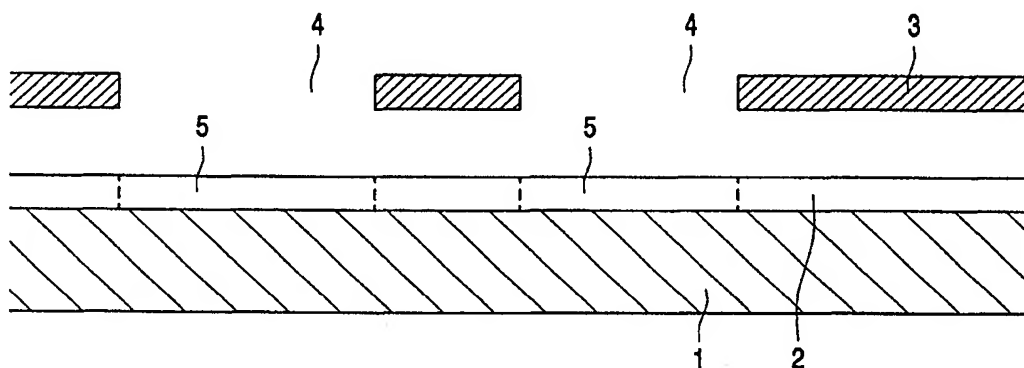




INTERNATIONAL APPLICATION PUBLISHED UNDER THE PATENT COOPERATION TREATY (PCT)

(51) International Patent Classification ⁷ : G02B 5/30, C09K 19/38	A1	(11) International Publication Number: WO 00/34808 (43) International Publication Date: 15 June 2000 (15.06.00)
(21) International Application Number: PCT/EP99/09711 (22) International Filing Date: 6 December 1999 (06.12.99) (30) Priority Data: 98204151.9 7 December 1998 (07.12.98) EP 99202635.1 13 August 1999 (13.08.99) EP (71) Applicant (for all designated States except US): KONINKLIJKE PHILIPS ELECTRONICS N.V. [NL/NL]; Groenewoudseweg 1, NL-5621 BA Eindhoven (NL). (72) Inventors; and (75) Inventors/Applicants (for US only): VAN DE WITTE, Peter [NL/NL]; Prof. Holstlaan 6, NL-5656 AA Eindhoven (NL). LUB, Johan [NL/NL]; Prof. Holstlaan 6, NL-5656 AA Eindhoven (NL). (74) Agent: BAELE, Ingrid, A., F., M.; Internationaal Octrooibureau B.V., Prof. Holstlaan 6, NL-5656 AA Eindhoven (NL).		(81) Designated States: CN, JP, KR, US, European patent (AT, BE, CH, CY, DE, DK, ES, FI, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE). Published <i>With international search report.</i> <i>Before the expiration of the time limit for amending the claims and to be republished in the event of the receipt of amendments.</i>

(54) Title: PATTERNED LAYER OF A POLYMER MATERIAL HAVING A CHOLESTERIC ORDER



(57) Abstract

The invention relates to a method of manufacturing a patterned layer of a polymer material having a cholesteric order, in which the material is oriented in such a way that the axis of the molecular helix of the cholesterically ordered material extends transversely to the layer, and in which the layer is patterned in that it comprises at least one region in which the pitch of the molecular helix differs from that of another region. This inventive method comprises the following steps: a) providing a layer of a cholesterically ordered material comprising a quantity of a convertible compound which in its non-converted and in its converted state determines the pitch of the cholesterically ordered material to a different extent, in which the conversion of said compound may be induced by radiation, b) irradiating the layer in accordance with a desired pattern so that at least a part of the convertible compound in the irradiated parts of the layer is converted, c) polymerizing and/or crosslinking the cholesterically ordered material to form a three-dimensional polymer. The convertible compound preferably comprises an isomerizable, chiral compound. Polymerization and/or crosslinking are preferably induced by irradiation using electron-beam radiation or actinic radiation.

FOR THE PURPOSES OF INFORMATION ONLY

Codes used to identify States party to the PCT on the front pages of pamphlets publishing international applications under the PCT.

AL	Albania	ES	Spain	LS	Lesotho	SI	Slovenia
AM	Armenia	FI	Finland	LT	Lithuania	SK	Slovakia
AT	Austria	FR	France	LU	Luxembourg	SN	Senegal
AU	Australia	GA	Gabon	LV	Latvia	SZ	Swaziland
AZ	Azerbaijan	GB	United Kingdom	MC	Monaco	TD	Chad
BA	Bosnia and Herzegovina	GE	Georgia	MD	Republic of Moldova	TG	Togo
BB	Barbados	GH	Ghana	MG	Madagascar	TJ	Tajikistan
BE	Belgium	GN	Guinea	MK	The former Yugoslav	TM	Turkmenistan
BF	Burkina Faso	GR	Greece		Republic of Macedonia	TR	Turkey
BG	Bulgaria	HU	Hungary	ML	Mali	TT	Trinidad and Tobago
BJ	Benin	IE	Ireland	MN	Mongolia	UA	Ukraine
BR	Brazil	IL	Israel	MR	Mauritania	UG	Uganda
BY	Belarus	IS	Iceland	MW	Malawi	US	United States of America
CA	Canada	IT	Italy	MX	Mexico	UZ	Uzbekistan
CF	Central African Republic	JP	Japan	NE	Niger	VN	Viet Nam
CG	Congo	KE	Kenya	NL	Netherlands	YU	Yugoslavia
CH	Switzerland	KG	Kyrgyzstan	NO	Norway	ZW	Zimbabwe
CI	Côte d'Ivoire	KP	Democratic People's	NZ	New Zealand		
CM	Cameroon		Republic of Korea	PL	Poland		
CN	China	KR	Republic of Korea	PT	Portugal		
CU	Cuba	KZ	Kazakstan	RO	Romania		
CZ	Czech Republic	LC	Saint Lucia	RU	Russian Federation		
DE	Germany	LI	Liechtenstein	SD	Sudan		
DK	Denmark	LK	Sri Lanka	SE	Sweden		
EE	Estonia	LR	Liberia	SG	Singapore		

Patterned layer of a polymer material having a cholesteric order.

5 The invention relates to a method of manufacturing a patterned layer of a cholesterically ordered polymer material, in which the axis of the molecular helix of the cholesterically ordered material extends transversely to the layer, and the layer is patterned in that it has at least an area in which the pitch of the molecular helix differs from that of another area.

The invention also relates to a layer of a cholesterically ordered polymer material.

The invention further relates to an information carrier provided with a patterned layer of a polymer material having a cholesteric order.

10 The invention further relates to a polarization conversion system provided with a patterned layer of a cholesterically ordered material.

The invention further relates to a cholesteric color filter having a patterned layer of a cholesterically ordered material.

15 The method of the type described in the opening paragraph is known per se. For example, United Kingdom patent specification GB 2,314,167 describes a patterned layer of a cholesterically ordered material. In accordance with this patent specification, such a layer may be manufactured by first providing a uniform layer of a cholesteric material on a substrate. By polymerizing areas of this layer at different temperatures, a patterned cholesteric layer is
20 obtained. Use is made of the fact that the pitch of the molecular helix of the cholesterically ordered material is temperature-dependent. By polymerizing areas of the layer at a given temperature, the pitch associated with this temperature is, as it were, frozen in these areas.

The known method has drawbacks. For example, in practice it has been found that the known method is difficult to implement. This notably applies to the case where more
25 than two areas having mutually different pitches must be provided in the layer. In that case, a relatively large number of masking steps is necessary and the precision with which the masks are adjusted is very critical. Moreover, the maximum difference in pitch which can be realized between the different areas by means of the known method appears to be relatively small. Patterning at different temperatures also appears to be difficult in practice.

CONFIRMATION COPY

It is an object of the invention to obviate the known drawbacks. More particularly, it is an object of the invention to provide a method in which the layer can be patterned at the same temperature and in which relatively large pitch differences between the different areas can be realized. It is a further object of the invention to provide a polarization
5 conversion system having a patterned layer of a cholesterically ordered material, and a cholesteric color filter having a patterned layer of a cholesterically ordered material, manufactured by means of this method.

These and other objects of the invention are achieved by means of a method of the type described in the opening paragraph, wherein the method comprises the steps of:

- 10 a. providing a layer of a cholesterically ordered material comprising a quantity of a convertible compound which in its non-converted and in its converted state determines the pitch of the cholesterically ordered material to a different extent, in which the conversion of said compound may be induced by radiation,
- b. irradiating the layer in accordance with a desired pattern so that at least a part of
15 the convertible compound in the irradiated parts of the layer is converted,
- c. polymerizing and/or crosslinking the cholesterically ordered material to form a three-dimensional polymer.

It has been found that, using the method according to the invention, that
20 patterned layers of cholesterically ordered, liquid crystalline material can be manufactured in a simple way at the same temperature, with the maximum pitch difference between the areas being relatively large. By (partially) converting the convertible compound in the irradiated areas of the layer, the pitch of the molecular helix in the layer is altered in these areas. The conversion of the convertible compound is effected by irradiation with energy in the form of,
25 for example, electro-magnetic radiation, nuclear radiation or an electron beam. Preferably said conversion is effected by means of UV radiation. The pitch of the molecular helix in the irradiated parts differs from the pitch of the molecular helix in the non-irradiated parts of the layer. By polymerizing and/or crosslinking the patterned layer thus obtained, the pitch in the different layer parts is frozen, as it were, and said pitch remains fixed during further process
30 steps, storage and use of the patterned layer. In this way, a patterned layer of cholesterically ordered material can be manufactured in a simple manner.

The degree of conversion of the convertible compound in a certain area is determined by the irradiation dose in said area. Consequently, the pitch of the molecular helix

is determined by the local irradiation dose. Said pitch of the molecular helix determines the local optical properties.

It is to be noted that, preferably, the cholesteric layer has a low absorbance for the radiation used in step b, and the radiation intensity along the axis of the helix (i.e. transverse to the layer) is relatively constant in each area. Consequently, the irradiation dose transverse to the layer is relatively constant, and therefore the value of the pitch, viewed along the axis of the helix, is relatively constant in each area. However, this value may differ for the different areas obtained by patterning. Viewed in the plane of the layer, the different areas are adjacent to each other, not subjacent.

When the cholesteric layer has a high absorbance for the radiation used in step b, the radiation intensity will show a gradient transverse to the layer according to Beer-Lambert's law. Consequently, the top of the layer will receive more radiation than the bottom of the layer. This will lead to the formation of a gradient in the pitch, viewed along the axis of the helix (i.e. transverse to the layer). The presence of an absorbing material in a non-absorbing cholesteric layer yields also a gradient in the pitch.

A variation in the pitch, transverse to the cholesteric layer, can be obtained by a method of manufacturing a layer of a cholesterically ordered polymer material, in which the material is oriented in such a way that the axis of the molecular helix of the cholesterically ordered material extends transversely to the layer, wherein the method comprises the following steps:

- a. providing a layer comprising a cholesterically ordered material, which material comprises a quantity of a convertible compound which in its non-converted and in its converted state determines the pitch of the cholesterically ordered material to a different extent, the conversion of said compound being inducible by radiation, and the layer substantially absorbs said radiation.
- b. irradiating the layer so that at least a part of the convertible compound in the irradiated parts of the layer is converted,
- c. polymerizing and/or crosslinking the cholesterically ordered material to form a three-dimensional polymer.

According to the prior art, a gradient in the pitch transverse to the layer could be obtained by a method described in the US patent US 5,793,456 (PHN 14.629), which discloses a method of manufacturing a cholesteric polarizer by providing a mixture of chiral

and nematogenic monomers, each having a different reactivity, in the form of a layer. The pitch of the molecular helix is governed to an important degree by the ratio between the chiral and the mesogenic monomer in the polymer material. Owing to the difference in reactivity between both monomers, the capture probability of the most reactive monomer is greater than that of the least reactive monomer. If during the polymerization of the mixture, which is initiated by actinic radiation, a variation in the radiation intensity is realized across the optically active layer to be formed, the most reactive monomer is preferably incorporated in the polymer at the locations of the highest radiation intensity. As a result, one or more concentration gradients of free monomer are formed during said polymerization process, causing monomer diffusion from locations with a low monomer concentration to locations with a high monomer concentration. This leads to an increase in reactive monomers in areas of the formed polymer material where, during the polymerization process, the radiation intensity was highest. As a result, the composition of the polymer material varies in the direction transverse to the polymer layer, causing a variation in the pitch of the molecular helix in the layer, which molecular helix is formed by the polymer. This variation of the pitch provides the optically active layer with a large bandwidth, the value of which is proportional to the value of the variation in pitch.

The known method of US 5,793,456 has drawbacks. For example, the speed of the process is governed by the diffusion of the monomer molecules which is an inherently slow process. The present invention provides a rapid method in which a pitch of the molecular helix can be made to vary transversely to the layer. An additional advantage of the method according to the invention is that the pitch of the cholesterically ordered material is determined by the irradiation dose instead of the irradiation intensity. A certain irradiation dose can be administered in a short timeframe using a high irradiation intensity.

A combination of the method according to claim 15 and the method described in US 5,793,456 is also possible. In said combination the effects of photo-isomerisation and diffusion are combined in order to obtain an even more increased gradient in the pitch of the molecular helix. If during the polymerization of a mixture of a photo-isomerisable chiral compound and a reactive nematogenic monomer, which polymerization is initiated by actinic radiation, a variation in the radiation intensity is realized across the optically active layer to be formed, two effects will cause a gradient in the pitch of the molecular helix. Firstly, due to the variation in the radiation intensity the amount of photo-isomerisable material that is converted will show the same variation across the thickness of the layer. Because the converted state determines the pitch of the cholesterically ordered material to a different extent, said variation

in converted material will yield a gradient in the pitch of the molecular helix. Secondly, the reactive nematogenic monomer is preferably incorporated in the polymer at the locations with the highest radiation intensity, as previously described in the third paragraph. As a result, the composition of the polymer material varies in the direction transverse to the polymer layer, causing a gradient in the pitch of the molecular helix, and at the end of the process, the cholesteric layer is crosslinked. When using the proper materials, the gradients of both effects add up to a substantial gradient in the pitch of the molecular helix, in a crosslinked layer, in a short time frame.

Preferably, the photo-isomerisable chiral compound is less reactive than the nematogenic monomer. In this sense, the photo-isomerisable chiral compound may be a chiral acrylate, or even a non-reactive compound, and the reactive nematogenic monomer may be a nematic diacrylate.

It has been found that, using the method according to claim 15, broadband polarizers of cholesterically ordered, liquid crystalline material can be manufactured in a simple way. By (partially) converting the convertible compound in the irradiated areas of the layer, the pitch of the molecular helix in the layer is altered in these areas. The conversion of the convertible compound is preferably effected by means of UV radiation. The degree of conversion of the convertible compound is determined by the irradiation dose. Consequently, the pitch of the molecular helix is determined by the local irradiation dose. Said pitch of the molecular helix determines the local optical properties. By polymerizing and/or crosslinking the layer thus obtained, the pitch in the different layer parts is frozen, as it were. In this way, a layer of cholesterically ordered material can be manufactured in a simple manner.

In contrast, when the cholesteric layer has a low absorbance for the radiation used in step b, the radiation intensity along the axis of the helix (i.e. transverse to the layer), is relatively constant over the cross-section of the layer. Consequently, the irradiation dose transverse to the layer is relatively constant, and therefore the value of the pitch, viewed along the axis of the helix, is relatively constant in each area, yielding a small bandwidth (approximately 60 – 70 nm).

An embodiment of the method according to claim 3 wherein the irradiation dose in accordance with step b decreases from the top to the bottom of the cholesteric layer, is

characterized in that, at the bottom of the cholesteric layer, said irradiation dose is less than 0.9 times the irradiation dose at the top of said layer.

When the cholesteric layer is caused to absorb the radiation used in accordance with step b, the radiation intensity will show a gradient transverse to the layer according to Beer-Lambert's law. Consequently, the top of the layer will receive more radiation than the bottom of the layer. Said variance in the irradiation dose over the cross-section of the layer will lead to the formation of a gradient in the pitch of the molecular helix, viewed along the axis of the helix (i.e. transverse to the layer). This gradient in the pitch provides the optically active layer with a larger bandwidth, the value of which is proportional to the value of the variation in the pitch. When the cholesteric material's absorbance of the radiation used in method step b is too small to produce the desired gradient in the pitch of the molecular helix in a certain amount of time, an absorbing material may be added to the cholesteric layer to obtain the required absorbance.

A preferred embodiment of the method according to claim 3 is characterized in that, at the bottom of the cholesteric layer, the irradiation dose in accordance with step b is less than 0.75 times the irradiation dose at the top of said layer. Said preferred variation of the irradiation dose across the thickness of the cholesteric layer yields a reflection band, which may span a substantial part of the visible spectrum.

From the description above, it is obvious that the pitch of the molecular helix in a layer of cholesterically ordered material can be suitably adjusted, yielding materials with special optical properties.

An interesting embodiment of the method according to the invention is characterized in that irradiation in accordance with step b is performed such that the irradiation dose is different for at least two different areas of the layer. By using this measure, it is possible to manufacture patterned cholesterically ordered layers which have juxtaposed areas, as viewed in the plane of the layer, with a different value of the pitch of the molecular helix. The different irradiation doses may be realized by using different irradiation periods at a substantially constant intensity. Alternatively, the different irradiation doses may be realized by using a higher irradiation intensity at a substantially constant irradiation period.

Preferably, however, masks are used having a number of apertures which have a different transmissivity to the radiation used. In that case, three or more areas in which the

pitch of the molecular helix is different can be obtained in one irradiation step, using one mask.

In principle, a large number of types of convertible compounds influencing the pitch of the molecular helix of cholesterically ordered material may be used within the scope of the invention. In the first place, convertible chiral compounds are feasible, which, due to irradiation, fall apart into non-chiral compounds. The presence of chiral compounds promotes the formation of a cholesteric ordering in a liquid crystalline solution. Irradiation of selected areas of a cholesterically ordered layer with decomposable chiral compounds leads to an increase of the pitch of the molecular helix in these areas.

Another advantageous embodiment of the method according to the invention is characterized in that the convertible compound comprises an isomerizable, chiral compound. The different isomeric forms of such an isomerizable chiral compound usually have a different influence on the pitch of the molecular helix of the cholesterically ordered material. By locally changing the ratio of these isomeric forms by way of irradiation, the pitch is changed. This provides an elegant possibility of manufacturing patterned layers of a polymer material with a cholesteric ordering and a different pitch. To prevent diffusion of the isomerizable, chiral compound in the patterned layer, this compound is preferably bound via a chemical bond to the liquid crystalline polymer material having the cholesteric order. In the latter case, the UV stability of the patterned layer also appears to have been improved.

The cholesterically ordered material used in the method according to the invention comprises liquid crystalline monomers, liquid crystalline oligomers and/or liquid crystalline linear polymers with reactive groups. Due to the presence of these reactive groups, this material can be converted into a polymer material by polymerization and/or into a three-dimensional molecular network by crosslinking. For the reactive groups, notably epoxy groups, vinyl ether groups and/or thiolene groups are suitable. Particularly suitable reactive groups are those of the (meth)acrylate type. It has been found that cholesterically ordered polymer layers having a high optical quality can be obtained with these types of reactive groups. It is to be noted that, when using linear polymers, only crosslinking is necessary for obtaining a three-dimensional network. However, when monomers and/or oligomers are used, polymerization and crosslinking should take place for obtaining the envisaged three-dimensional molecular network. The stabilizing of the cholesteric layer in process step c after selective adjustment of the pitch of the cholesterically ordered layer in process step b, is an important step in the method according to the invention.

An embodiment of the method according to the invention is therefore characterized in that the polymerization and/or crosslinking is initialized and/or catalyzed by the addition of an initiator or catalyst from the fluid or gaseous phase. Said addition is preferably performed after steps a and b in accordance with the invention in order to prevent a polymerization and/or crosslinking reaction during steps a and b. Various initiators and catalysts are applicable.

A further embodiment of the method according to the invention is therefore characterized in that polymerization and/or crosslinking is induced by a thermally decomposable initiator. In that case, the layer of cholesterically ordered material preferably comprises a small quantity of a thermally decomposable polymerization initiator. Said initiator is inactive during process step b according to the invention. Subsequently, the polymerization and/or crosslinking of process step c, may be effected by activating the initiator at an elevated temperature.

An embodiment of the method according to the invention is characterized in that polymerization and/or crosslinking is effected by means of electron-beam irradiation. Very hard layers can be manufactured by means of this method. In this variant of the method according to the invention, it is not necessary to use a polymerization initiator.

An embodiment of the method according to the invention is characterized in that polymerization and/or crosslinking is effected by exposure to actinic radiation. The polymerization and/or crosslinking of a layer of the cholesterically ordered material (step c) can take place in the presence of a photo-initiator by using actinic radiation such as UV radiation. An advantage of using photo-polymerization is that this method permits local polymerization and/or crosslinking in very small areas.

Since the conversion of the convertible compound (step b) is also preferably effected by means of UV radiation, step b and step c of the method claimed may interfere with one another. In order to disentangle these method steps, the next three preferred embodiments of the method, as described below, may be used:

A first preferred embodiment of the method according to the invention is characterized in that the irradiation in accordance with step b is performed at a temperature, at

which the polymerization and/or crosslinking reaction is substantially hampered. The polymerization and/or crosslinking reaction rate is temperature-dependent. At a low temperature (at a high viscosity), the polymerization and/or crosslinking reaction is slower than the reorientation of the cholesteric helix, thus allowing the pitch of the cholesterically ordered material to be adjusted with limited or substantially no polymerization and/or crosslinking. At a high temperature (at a low viscosity), the polymerization and/or crosslinking reaction is faster than the reorientation of the cholesteric helix, thus allowing a polymerization and/or crosslinking with a limited or substantially no change in the pitch of the cholesterically ordered material. In conclusion, applying two irradiation steps, each at a different temperature, disentangles method steps b and c.

A second preferred embodiment of the method according to the invention is characterized in that the irradiation in accordance with step b is performed by means of radiation which is substantially inert with respect to the initiation of the polymerization and/or crosslinking reaction. This method uses two irradiation steps, each step using radiation with a different wavelength. According to this embodiment of the method according to the invention, the layer is irradiated in step b with a desired pattern using radiation which is substantially inert with respect to the initiation of the polymerization and/or crosslinking reaction, thus allowing the pitch of the cholesterically ordered material to be adjusted with limited or substantially no polymerization and/or crosslinking. Subsequently, the polymerization and/or crosslinking is effected by means of actinic radiation having a suitable wavelength to initiate polymerization and/or crosslinking. Preferably, but not necessarily, said actinic radiation has a wavelength with a limited or substantially no influence on the pitch of the cholesterically ordered material. If said actinic radiation induces an additional change in the pitch of the cholesterically ordered material, then this must be taken into account when setting the pitch of the cholesterically ordered material in process step b.

A third preferred embodiment of the method according to the invention is characterized in that the irradiation in accordance with step b is performed in an atmosphere which substantially hampers the polymerization and/or crosslinking reaction. This method uses two irradiation steps, each step being performed in a different atmosphere. According to this embodiment of the method according to the invention, the layer is irradiated in step b with a desired pattern in an atmosphere comprising molecules, e.g. oxygen or nitrogen-monoxide, that quench the activated photo-initiators. Said quenching essentially deactivates the activated photo-initiators and thereby hampers the polymerization and/or crosslinking reaction. Consequently, the pitch of the cholesterically ordered material can be adjusted with limited or

substantially no polymerization and/or crosslinking. Subsequently, the polymerization and/or crosslinking in accordance with step c is initiated by irradiation in a non-quenching atmosphere, e.g. nitrogen. Preferably, but not necessarily, the second wavelength used for the irradiation in accordance with step c, has a limited or substantially no influence on the pitch of the cholesterically ordered material. If said actinic radiation induces an additional change in the pitch of the cholesterically ordered material, then this must be taken into account when setting the pitch of the cholesterically ordered material in process step b.

The method according to the invention may yield a patterned layer of a polymer material having very interesting optical properties due to the selectively adjusted cholesteric ordering. Said method is particularly suitable for the manufacture of information carriers which may comprise (high resolution) data, text, images, emblems, logos, holograms or gratings.

The invention relates to a layer of a cholesterically ordered polymer material, in which the material is oriented in such a way that the axis of the molecular helix of the cholesterically ordered material extends transversely to the layer, which layer comprises a quantity of a convertible compound which in its non-converted and in its converted state determines the pitch of the cholesterically ordered material to a different extent, and which layer comprises at least one region in which the pitch of the molecular helix differs from that of another region. This layer may be manufactured by means of one of the methods as described hereinbefore.

The invention also relates to an information carrier provided with a patterned layer of a polymer material having a cholesteric order, manufactured by means of one of the methods as described hereinbefore.

The invention further relates to a polarization conversion system provided with a patterned layer of a polymer material having a cholesteric order, manufactured by means of one of the methods as described hereinbefore.

The invention further relates to a color filter with a patterned layer of a polymer material having a cholesteric order, manufactured by means of one of the methods as described hereinbefore.

The invention further relates to a layer of a cholesterically ordered polymer material, in which the material is oriented in such a way that the axis of the molecular helix of the cholesterically ordered material extends transversely to the layer, and the pitch of the

molecular helix varies transversely to the layer, said layer being manufactured by means of one of the methods as described hereinbefore.

The invention further relates to a polarizer comprising a layer having a cholesteric order as described in the previous paragraph, which layer is manufactured by means of one of the methods as described hereinbefore.

These and other aspects of the invention are apparent from and will be elucidated with reference to the embodiments described hereinafter.

In the drawings:

Fig. 1 diagrammatically shows some process steps of the method according to the invention,

Fig. 2 shows the chemical structural formulas of some compounds used in the methods according to the invention,

Fig. 2A shows the monomer RM82 (supplied by Merck),

Fig. 2B shows the E isomer of the isomerizable chiral compound (E) – (1R, 4R) – 2 – (4 – (4 – hexyloxybenzoyloxy) phenyl – methene) menthan – 3 – on, which is a reactive menthone (RM),

Fig. 2C shows the monomer RM257 (supplied by Merck),

Fig. 2D shows the molecule of the material referred to as TetraCN.

Fig. 2E shows the molecule of the material referred to as EtherLC.

Fig. 2F shows a copolymer having a certain amount of component A as side chains. The copolymer will be referred to as M15, whereby the number represents the mol % of menthone units.

Fig. 3 is a diagrammatic cross-sectional view of a polarization conversion system provided with a patterned layer of a polymer material having a cholesteric order, manufactured by means of a method according to the invention,

Fig. 4 is a diagrammatic plan view of a color filter with a patterned layer of a polymer material having a cholesteric order, manufactured by means of the method according to the invention,

Fig. 5 illustrates the broadening of the reflection band as a function of the irradiation time, and

Fig. 6 shows the reflection spectra of a broadband polarizer.

It is to be noted that, for the sake of clarity, the Figures are not drawn to scale.

The essence of the method according to the invention will be elucidated with reference to Figure 1. This Figure shows a substrate 1 of, for example glass, a main surface of which is provided with an orientation layer (not shown) of, for example, polyimide. A layer of a liquid crystalline material 2 having a thickness of approximately 2 μm is provided on this layer by means of a thick-film technique (doctor blade coating or spin-coating). This layer is given a cholesteric order by means of the orientation layer. The material is oriented in such a way that the molecular helix of the cholesterically ordered material extends transversely to the layer.

In the relevant case, the liquid crystalline material consisted of a mixture of compound A (RM82, Merck) to which 0.5% by weight of the isomerizable, chiral compound B (RM) had been added. The chemical structural formulas of these compounds are shown in Fig. 2. The compound B induces the formation of the molecular helix in the liquid crystalline material so that the desired cholesteric order is obtained. In the case described, the helix had a pitch of 7.5 μm .

Subsequently a mask 3 with apertures 4 was positioned above the layer 2 of the cholesterically ordered liquid crystalline material. Layer 2 was selectively irradiated at ambient temperature for 400 sec using UV radiation (365 nm, 8 mW/cm²). Due to this irradiation, a large part of the isomerizable, chiral compound B in the irradiated parts 5 was converted from the E form to the Z form. These two forms of the chiral compound have a different influence on the helix of the cholesterically ordered material. It has been found that the pitch of the irradiated parts 5 increased from 7.5 to 40 μm .

After the irradiation, the mask was removed and the layer was irradiated with an electron-beam. As a result, polymerization and crosslinking of the liquid crystalline material took place. During this treatment, the chiral compound was also polymerized so that it was incorporated in the three-dimensional structure of the cholesterically ordered, polymerized material. The patterned layer 2 thus formed was subsequently removed from the substrate 1.

The polymerization and/or crosslinking step may alternatively be accomplished by means of actinic radiation, as the next examples illustrate.

Example 1.

A mixture is made which is composed of 42.5 % of a vitrifying liquid crystal compound D (TetraCN, $T_g = 80\text{ }^{\circ}\text{C}$), 13 % of compound B (RM), 42.5 % of a vitrifying liquid crystal compound E having two acrylic moieties (EtherLC, $T_g = -20\text{ }^{\circ}\text{C}$), and 2 %
5 photoinitiator (Irgacure 651, Ciba Specialty Chemicals). This combination of compound D and E yields a reactive mixture, which is vitrified at room temperature. Component B is a photo-isomerizable component, which is used to set the pitch of the cholesterically ordered layer. The photo-initiator initiates the polymerization and/or crosslinking reaction.

Mixing is performed at $120\text{ }^{\circ}\text{C}$, yielding a mixture which reflects green light.

10 Said mixture is deposited as a thin film (5 micrometers thick) between two substrates (e.g. glass plates) provided with an orientation layer adjacent to the mixture. Subsequently, this cholesterically ordered film is cooled down to room temperature, and irradiated with UV light (365 nm) for a short period of time, according to a suitable pattern. Surprisingly, it seems as if the irradiation at room temperature has no effect on the cholesteric film. After irradiation, the
15 temperature of the film was raised to $120\text{ }^{\circ}\text{C}$. During this heating, however, the color of the irradiated areas still underwent a change. The non-irradiated areas did not change color at all. Being at $120\text{ }^{\circ}\text{C}$, the cholesteric film is substantially liquid. The polymerisation and/or crosslinking is subsequently initiated by irradiating with UV light (365 nm) for a short period of time according to a suitable pattern. Immediately after this irradiation at high temperature,
20 the irradiated areas solidify due to polymerization and/or crosslinking. The non-irradiated areas remained in a substantially liquid form.

This example also demonstrates that actinic radiation, such as UV light, can be used for both method steps b and c, and that applying two irradiation steps, each at a different temperature, disentangles method steps b and c.

25

Example 2.

In another example, a mixture is made of 15 % of compound B (RM), 84 % of compound C (RM257), and 1 % Darocure 4265. Darocure 4265 is a photo-initiator which can be activated using 405 nm UV radiation. Said mixture is deposited as a thin film, and
30 subsequently, at room temperature and under ambient atmosphere (air, containing 20 % oxygen), irradiated with UV light (365 nm) for a short period of time, according to a suitable pattern. During this irradiation the pitch of the molecular helix in the layer of cholesterically ordered material is altered, and the photo-initiator is activated. However, the polymerization and crosslinking of said layer is hampered because the activated initiator is quenched by the

oxygen in the ambient atmosphere. After setting the required pitch, the layer is irradiated with 405 nm UV light under a nitrogen atmosphere. During this irradiation the layer is polymerized and crosslinked with substantially no change in the pitch of the molecular helix in the layer.

5 Figure 3 is a cross-sectional view of a polarization conversion system. This system comprises an optically transparent body 11 of, for example glass which consists mainly of two parts 12 and 13. These parts are separated by a polarization-selective layer 14. A reflecting mirror 15 which extends substantially parallel to the polarization-selective layer 14 is provided on a surface of one of the two parts.

10 The exit plane 16 of the system is also provided with a patterned layer 17 of a polymer material having a cholesteric order. This layer comprises two sub-layers 18 and 19. The thickness and cholesteric order of sub-layer 18 are chosen to be such that the direction of polarization of an ongoing, polarized light beam having the desired wavelength is rotated exactly 90°. The cholesteric order of sub-layer 19 is adapted in such a way that the pitch has a
15 substantially infinite value. When a polarized light beam passes through sub-layer 19, the direction of polarization of this beam is not rotated in such a case.

 An unpolarized light beam p, s which incidents on the entrance plane 20 of the polarization system, is split on the polarization-selective layer 14 into an ongoing polarized light beam p and a reflected polarized light beam s. The ongoing light beam p leaves the
20 system via exit plane 16 and sub-layer 19. The direction of polarization of this light beam p is not changed when it passes through sub-layer 19. The reflected light beam s is reflected at mirror 15 in such a way that this light beam, which is parallel to light beam p, leaves the system via exit plane 16 and sub-layer 18. The direction of polarization of light beam s is rotated 90° in sub-layer 18 so that a light beam p is obtained. The two exiting light beams thus
25 have the same direction of polarization.

 The patterned layer is essentially manufactured in the same way as described with reference to Fig. 1. However, in this case, a synthetic material substrate was used, whose surface dimensions corresponded to the dimensions of the exit plane of the polarization conversion system. The substrate was provided with a 2.5 µm thick layer of a nematic liquid
30 crystalline compound C to which 0.6% by weight of the isomerizable, chiral compound B (E-isomer) had been added, as well as 0.3% by weight of the non-photoisomerizable compound R811 (Merck, structural formula not shown). The addition of the non-photoisomerizable chiral dopant R811, having an opposite twisting sense with respect to compound B, is required to compensate for the residual twisting power of the Z-isomer of compound B.

After the material had been cholesterically ordered, the pitch of the molecular helix was defined at 10 μm . In combination with a layer thickness of approximately 2.5 μm , this leads to a rotation of approximately 90° of the polarization for visible light. Subsequently, half the layer was irradiated with UV light (365 nm, 20 mW/cm²) for a long period of time via a mask, so that all E isomer of compound B was converted into Z isomer. As a result, the pitch of the cholesterically ordered material became approximately infinite in the irradiated part of the layer, so that, in fact, a nematically ordered material was obtained. The direction of polarization of polarized visible light is not rotated when this light passes through this layer. After polymerization and crosslinking, the layer was removed from the substrate and provided on the exit plane of the system.

It is apparent that patterned polarization rotators can be made by means of the method as described hereinbefore, where the local twist angle of the cholesterically ordered layer and thereby the local rotation of the polarization of a transmitted light beam, is determined by the dose of UV light at that particular position. Lateral resolutions smaller than 10 micron can easily be obtained. Local control of the polarization rotation offers new possibilities in the preparation of optical components. Potential applications are, for example, patterned polarization rotators (as described above), optical data storage media, polarization sensitive gratings, polarization sensitive recordings, or holograms for copy protection.

Moreover, between two polarizers, variations in polarization rotation show up as variations in light transmission. Full grey-scale capability is achieved by adjusting the twist angle in a range of 90 degrees. Preferably, the twist angle is adjusted in the region around 0 degrees, because in this region the polarization rotation has a low wavelength dependence.

Fig. 4 is a plan view of a color filter which mainly consists of a 5 μm thick, patterned layer of a polymer material having a cholesteric order. This layer comprises an array of three sub-layers R (red), G (green) and B (blue) which are regularly spread across the layer. The sub-layers had dimensions of 100 μm x 100 μm . The pitch of the molecular helix of the cholesterically ordered material in the different sub-layers was 0.28 μm (blue reflection), 0.34 μm (green reflection) and 0.41 μm (red reflection), respectively.

The color filter was manufactured as follows. A substrate was provided with an approximately 5 μm thin layer of compound C in which 15% by weight of the compound B was incorporated. After ordering, the entire layer had a pitch of the molecular helix of

0.28 μm , which yields the desired blue reflection B. Subsequently, two thirds of the layer were irradiated (365 nm, 5.6 mW/cm²) according to a suitable pattern for 90 seconds so that the irradiated areas had a pitch of the molecular helix of 0.34 μm , which yields the desired green reflection G. Subsequently, half of the areas of the layer that had already been irradiated were
5 further irradiated for 90 seconds so that these twice irradiated areas had a pitch of the molecular helix of 0.41 μm , which yields the desired red reflection R. Finally, the complete layer was polymerized and crosslinked by means of electron-beam irradiation. A color filter array suitable for use in a liquid crystal display (LCD) was obtained.

In an alternative preferred embodiment of the method according to the
10 invention of manufacturing the above-mentioned color filter array, a grey-tint mask was used. In this mask, the transmission of the mask apertures intended for the red pixels is 100 %, the transmission of the mask apertures intended for the green pixels is 50%, and the UV radiation is completely blocked at the positions intended for the blue pixels. The three envisaged areas having a different pitch can be obtained with this mask in a single irradiation step of 180 sec.
15 Also in this case, the separate pixels had dimensions of 100 μm x 100 μm .

The method according to the invention has proved that it is possible to manufacture a patterned layer of a polymer material having a cholesteric order, in which the layer has juxtaposed areas where the pitch of the molecular helix differs from that of another
20 area. This method has proved that it is possible to manufacture patterned cholesteric layers for use in a polarization conversion system and a color filter.

However, the method according to the invention may also lead to a layer of a cholesterically ordered polymer material, in which the material is oriented in such a way that
25 the axis of the molecular helix of the cholesterically ordered material extends transversely to the layer, and the pitch of the molecular helix is made to vary transversely to the layer. The method to obtain such a layer, as claimed in claim 3, will be elucidated with reference to Figure 1. This Figure shows a substrate 1 of, for example, glass, a main surface of which is provided with an orientation layer (not shown) of, for example, polyimide. A layer of a liquid
30 crystalline material 2 having a thickness of approximately 10 μm is provided on this layer by means of a thick-film technique (doctor blade coating or spin-coating). This layer is given a cholesteric order by means of the orientation layer. The material is oriented in such a way that the molecular helix of the cholesterically ordered material extends transversely to the layer.

In the relevant case, the liquid crystalline material consisted of a mixture of 98 % by weight of M15, a copolymer with 15 mol % of menthone units (compound 2F), and 2 % by weight of an UV absorber (tinuvin 1130, Ciba specialty Chemicals). The chemical structural formulas of some of these compounds are shown in Fig. 2. The copolymer induces the formation of the molecular helix in the liquid crystalline material so that the desired cholesteric order is obtained. In the case described, the helix had a pitch of approximately 0.3 μm .

Subsequently layer 2 was irradiated at ambient temperature in air (containing the normal amount of oxygen) for 10 minutes using UV radiation (365 nm, 2.8 mW/cm²). Due to this irradiation, a part of the isomerizable, chiral compound B in the irradiated layer was converted from the E form to the Z form. Consequently, the color of the layer shifts towards longer wavelengths (red shift).

Due to the presence of the UV absorber, the intensity of the radiation in the cholesteric layer shows a gradient transverse to the layer. The top of the sample receives more UV light than the bottom of the sample, resulting in a gradient in the amount of converted isomerizable compound. This gradient yields a corresponding gradient in the pitch of the helix of the cholesteric layer. Consequently, the red shift at the bottom of the layer is lower than the red shift at the top of the layer, causing a broadening of the reflection band as demonstrated in figure 5.

After establishing the required reflection band, the layer was irradiated with an electron-beam in a nitrogen atmosphere. As a result, crosslinking of the liquid crystalline material took place.

The polymerization and/or crosslinking step may alternatively be accomplished by means of actinic radiation when a small amount of a photo-initiator is added to the mixture of liquid crystalline material. If the photo-initiator is activated by light having the same wavelength as that used for process step b, then step b and step c can be performed simultaneously.

For example, a liquid crystalline material consisted of a mixture of 15 % by weight of menthone-acrylate (compound 2A), 82 % by weight of RM257 (compound 2C), 2 % by weight of an UV absorber (tinuvin 1130, Ciba Specialty Chemicals), 1 % by weight of a photo-initiator (Darocure 4265, Ciba Specialty Chemicals), and 300 ppm of an inhibitor (methoxyphenol). The chemical structural formulas of some of these compounds are shown in Figure 2. A layer of this mixture is provided on a substrate, which comprises an orientation

layer of, for example, polyimide. This layer is given a cholesteric order by means of the orientation layer. The material is oriented in such a way that the molecular helix of the cholesterically ordered material extends transversely to the layer.

Subsequently, the layer was irradiated for 3 minutes using UV radiation (365 nm, 6 mW/cm²). Due to this irradiation, a part of the isomerizable, chiral compound A in the irradiated layer was converted from the E form to the Z form. Consequently, the color of the layer shifts towards longer wavelengths (red shift). Due to the presence of the UV absorber, the intensity of the radiation in the cholesteric layer shows a gradient transverse to the layer, resulting in a gradient in the pitch of the helix of the cholesteric layer, causing a broadening of the reflection band. Due to the photo-initiator, the polymerization and crosslinking reaction is initiated. This reaction is hampered by an inhibitor in order to allow the pitch and the gradient in the pitch of the helix of the cholesteric layer to be developed before the three-dimensional network is formed.

When the polymerization and/or crosslinking step is accomplished by means of actinic radiation, and process steps b and c should be disentangled, process step b may be performed in an oxygen atmosphere which hampers the polymerization and crosslinking if the selected activated initiator is quenched by the oxygen in the ambient atmosphere. For initiating said crosslinking, the irradiation process in accordance with step c is performed in a nitrogen atmosphere.

In another example, a mixture is made of 15 % by weight of menthone-acrylate (compound 2A), 82 % by weight of RM257 (compound 2C), 2 % by weight of an UV absorber (tinuvin 1130, Ciba Specialty Chemicals), 1 % by weight of a photo-initiator (Darocure 4265, Ciba Specialty Chemicals). Said mixture is deposited as a thin film on a substrate, and subsequently irradiated with UV light (365 nm, 1 mW/cm²) during 5 minutes. The irradiation is performed under ambient atmosphere (air, containing the usual amount of oxygen). Due to this irradiation, a part of the isomerizable, chiral compound A in the irradiated layer was converted from the E form to the Z form. Consequently, the color of the layer shifts towards longer wavelengths (red shift). Due to the presence of the UV absorber, the intensity of the radiation in the cholesteric layer shows a gradient transverse to the layer, resulting in a gradient in the pitch of the helix of the cholesteric layer, causing a broadening of the reflection band. Due to the photo-initiator, the polymerization and crosslinking reaction is initiated. This reaction is hampered by the oxygen in the ambient atmosphere.

When the desired pitch and the gradient in the pitch of the helix of the cholesteric layer is established, the layer is polymerized and crosslinked using 405 nm UV light (4 mW/cm^2) in a nitrogen atmosphere. At this wavelength, the photo-initiator Darocure 4265 can be activated without substantial conversion of the isomerizable compound A. During
5 this irradiation the layer is polymerized and crosslinked with substantially no change in the pitch of the molecular helix in the layer.

It is apparent that a broadband polarizer can be made by means of the method as described hereinbefore. Said broadband polarizer comprises a layer of a polymer material
10 having a cholesteric order, the material being oriented in such a manner that the axis of the molecular helix extends transversely to the layer, and the pitch of the molecular helix in the layer is varied

In general, the invention relates to a method of manufacturing a patterned layer
15 of a polymer material having a cholesteric order, in which the material is oriented in such a way that the axis of the molecular helix of the cholesterically ordered material extends transversely to the layer, and in which the layer comprises at least one region in which the pitch of the molecular helix differs from that of another region. This inventive method comprises the following steps:

- 20 a. providing a layer of a cholesterically ordered material comprising a quantity of a convertible compound which in its non-converted and in its converted state determines the pitch of the cholesterically ordered material to a different extent, in which the conversion of said compound may be induced by radiation,
- b. irradiating the layer in accordance with a desired pattern so that at least a part of
25 the convertible compound in the irradiated parts of the layer is converted,
- c. polymerizing and/or crosslinking the cholesterically ordered material to form a three-dimensional polymer.

The convertible compound preferably comprises an isomerizable, chiral compound. Polymerization and/or crosslinking are preferably induced by irradiation using
30 electron-beam radiation or actinic radiation.

CLAIMS:

1. A method of manufacturing a patterned layer of a polymer material having a cholesteric order, in which the material is oriented in such a way that the axis of the molecular helix of the cholesterically ordered material extends transversely to the layer, and the layer is patterned in that it has least an area in which the pitch of the molecular helix differs from that of another area, wherein the method comprises the steps of:

- a. providing a layer of a cholesterically ordered material comprising a quantity of a convertible compound which in its non-converted and in its converted state determines the pitch of the cholesterically ordered material to a different extent, in which the conversion of said compound may be induced by radiation,
- 10 b. irradiating the layer in accordance with a desired pattern so that at least a part of the convertible compound in the irradiated parts of the layer is converted,
- c. polymerizing and/or crosslinking the cholesterically ordered material to form a three-dimensional polymer.

15 2. A method as claimed in claim 1, characterized in that irradiation in accordance with step b is performed such that the irradiation dose is different for at least two different areas of the layer.

20 3. A method of manufacturing a layer of a cholesterically ordered polymer material, in which the material is oriented in such a way that the axis of the molecular helix of the cholesterically ordered material extends transversely to the layer, wherein the method comprises the following steps:

- a. providing a layer comprising a cholesterically ordered material, which material comprises a quantity of a convertible compound which in its non-converted and in its converted state determines the pitch of the cholesterically ordered material to a different extent, the conversion of said compound being inducible by radiation, and the layer substantially absorbs said radiation,
- 25 b. irradiating the layer so that at least a part of the convertible compound in the irradiated parts of the layer is converted,

c. polymerizing and/or crosslinking the cholesterically ordered material to form a three-dimensional polymer.

4. A method as claimed in claim 3, wherein the irradiation dose in accordance with step b decreases from the top to the bottom of the cholesteric layer, is characterized in that, at the bottom of the cholesteric layer, said irradiation dose is less than 0.9 times the irradiation dose at the top of said layer.

5. A method as claimed in claim 3, characterized in that, at the bottom of the cholesteric layer, the irradiation dose in accordance with step b is less than 0.75 times the irradiation dose at the top of said layer.

6. A method as claimed in claim 1 or 3, characterized in that irradiation in accordance with step b is performed via a mask having a number of apertures which have different transmissivity to the radiation used.

7. A method as claimed in claim 1 or 3, characterized in that the convertible compound comprises an isomerizable, chiral compound.

8. A method as claimed in claim 1 or 3, characterized in that the polymerization and/or crosslinking is initialized and/or catalyzed by the addition of an initiator or catalyst from the fluid or gaseous phase.

9. A method as claimed in claim 1 or 3, characterized in that the polymerization and/or crosslinking is induced by a thermally decomposable initiator.

10. A method as claimed in claim 1 or 3, characterized in that polymerization and/or crosslinking is effected by means of electron-beam irradiation.

11. A method as claimed in claim 1 or 3, characterized in that polymerization and/or crosslinking is effected by exposure to actinic radiation.

12. A method as claimed in claim 11, characterized in that irradiation in accordance with step b is performed at a temperature at which the polymerization and/or crosslinking reaction is substantially hampered.
- 5 13. A method as claimed in claim 11, characterized in that irradiation in accordance with step b is performed by means of radiation which is substantially inert with respect to the initiation of the polymerization and/or crosslinking reaction.
- 10 14. A method as claimed in claim 11, characterized in that irradiation in accordance with step b is performed in an atmosphere which substantially hampers the polymerization and/or crosslinking reaction
- 15 15. A layer of a cholesterically ordered polymer material, in which the material is oriented in such a way that the axis of the molecular helix of the cholesterically ordered material extends transversely to the layer, which layer comprises a quantity of a convertible compound which in its non-converted and in its converted state determines the pitch of the cholesterically ordered material to a different extent, and which layer comprises at least one region in which the pitch of the molecular helix differs from that of another region.
- 20 16. An information carrier comprising a patterned layer of a polymer material having a cholesteric order, as claimed in claim 15.
- 25 17. A polarization conversion system comprising a patterned layer of a polymer material having a cholesteric order, as claimed in claim 15.
18. A color filter comprising a patterned layer of a polymer material having a cholesteric order, as claimed in claim 15.
- 30 19. A layer of a cholesterically ordered polymer material as claimed in claim 15, in which the material is oriented in such a way that the axis of the molecular helix of the cholesterically ordered material extends transversely to the layer, and the pitch of the molecular helix varies transversely to the layer.

20. A polarizer comprising a layer of a polymer material having a cholesteric order, as claimed in claim 19.

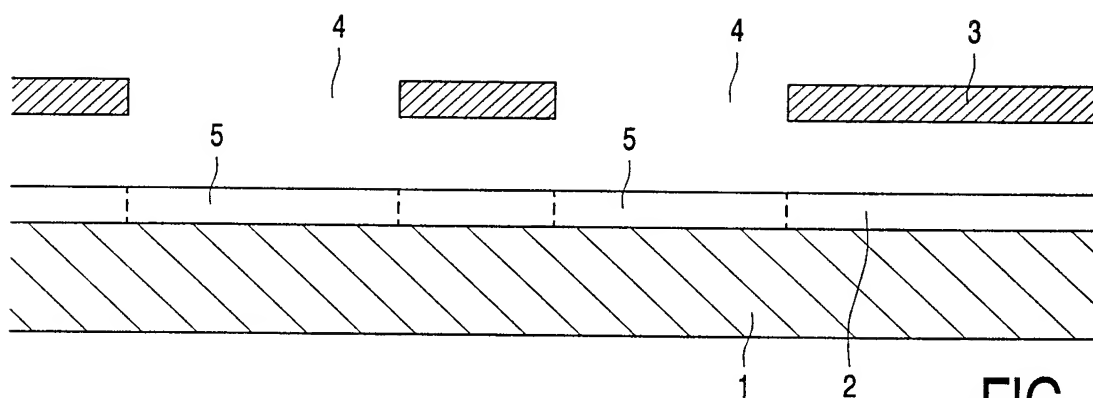
$\frac{1}{5}$ 

FIG. 1

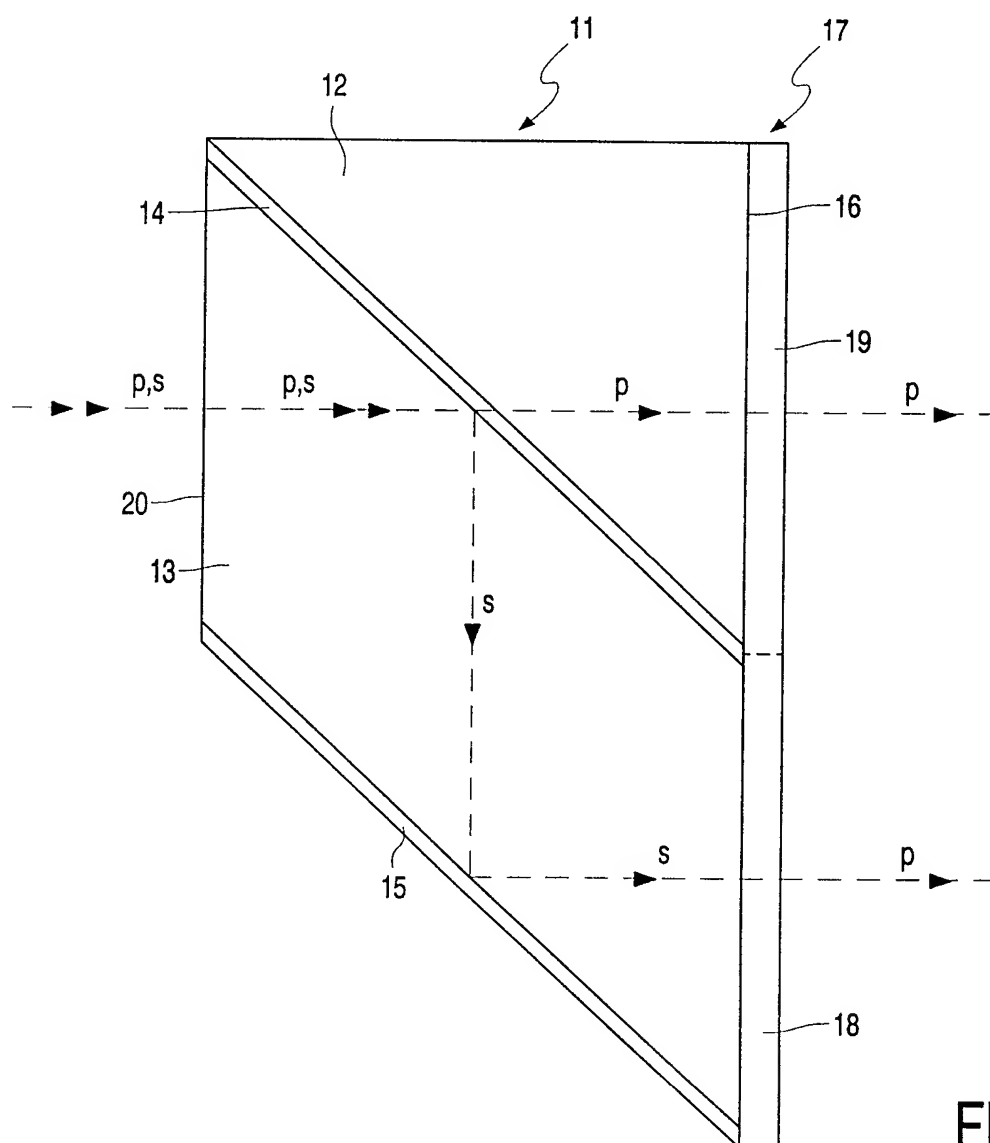


FIG. 3

2/5

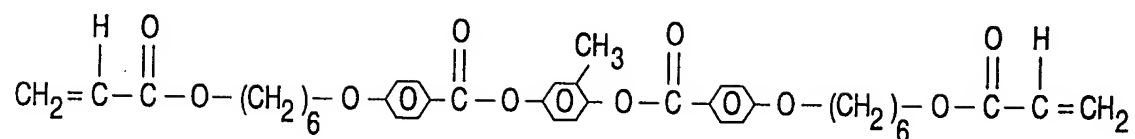


FIG. 2A

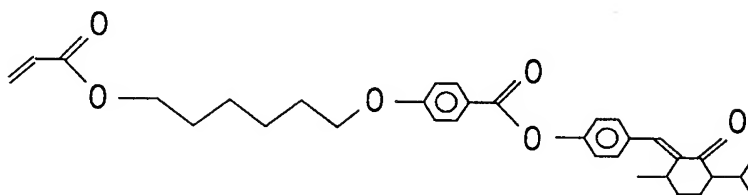


FIG. 2B

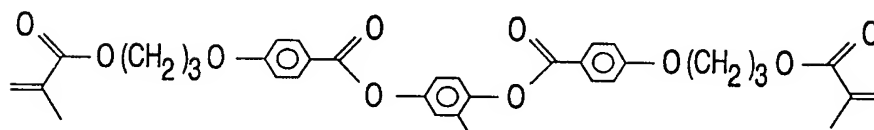


FIG. 2C

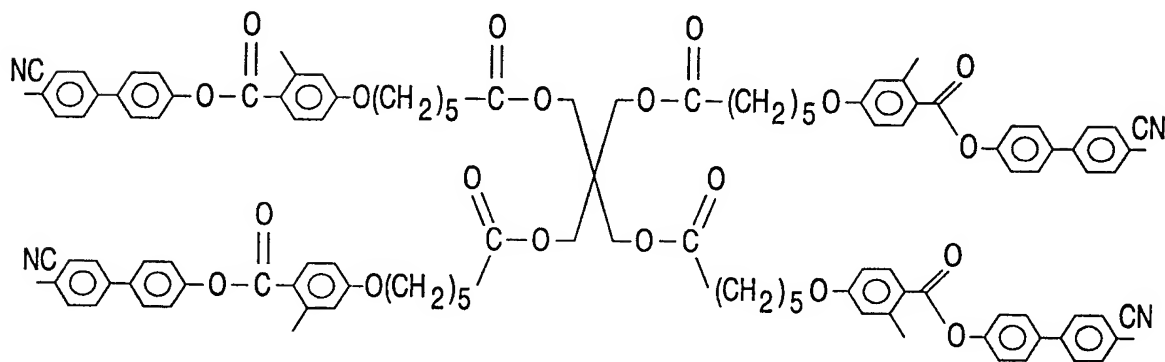


FIG. 2D

3/5

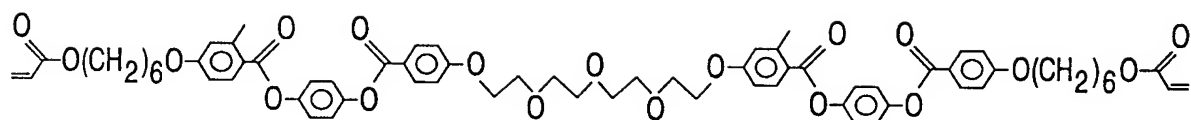


FIG. 2E

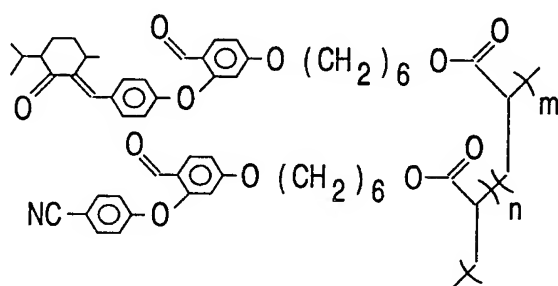


FIG. 2F

4/5

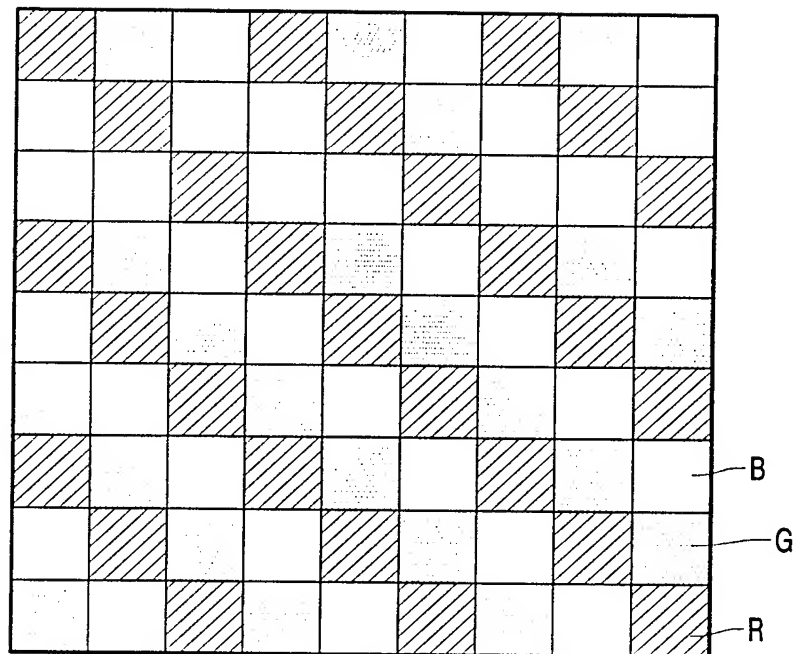


FIG. 4

5/5

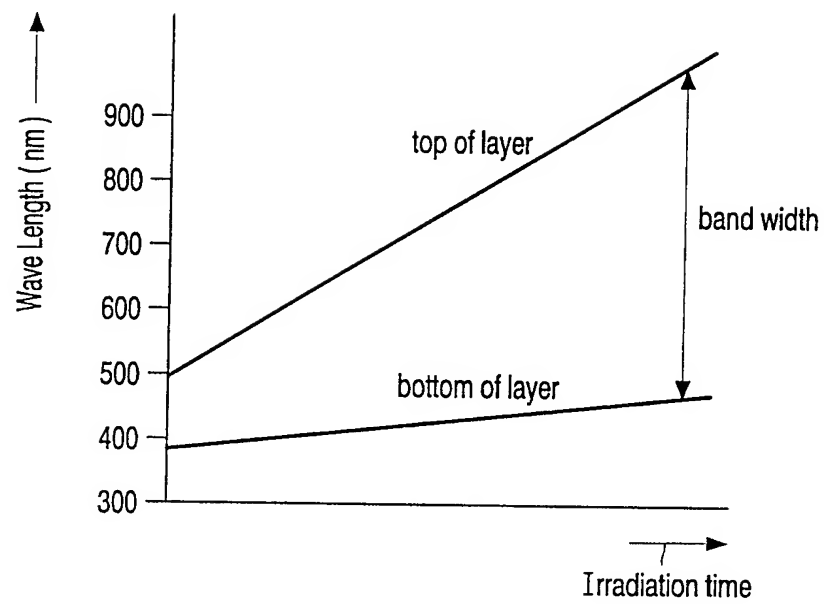


FIG. 5

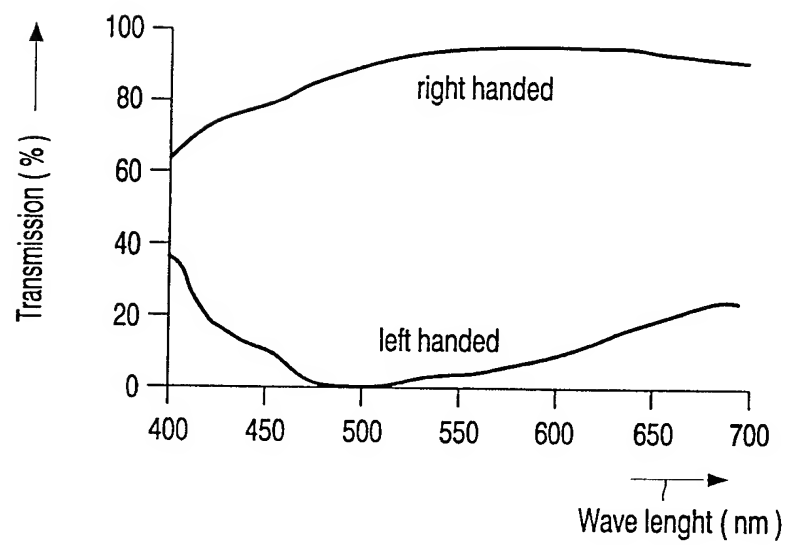


FIG. 6

INTERNATIONAL SEARCH REPORT

International Application No.

PCT/EP 99/09711

A. CLASSIFICATION OF SUBJECT MATTER

IPC 7 G02B5/30 C09K19/38

According to International Patent Classification (IPC) or to both national classification and IPC

B. FIELDS SEARCHED

Minimum documentation searched (classification system followed by classification symbols)

IPC 7 G02B C09K

Documentation searched other than minimum documentation to the extent that such documents are included in the fields searched

Electronic data base consulted during the international search (name of data base and, where practical, search terms used)

C. DOCUMENTS CONSIDERED TO BE RELEVANT

Category *	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
X	US 5 668 614 A (CHIEN LIANG-CHY ET AL) 16 September 1997 (1997-09-16) abstract column 2, line 26 - line 49 column 3, line 22 - line 32 column 4, line 1 - line 10 column 37, line 18 - line 27; claim 23	1-3, 7, 15-18
A	US 5 793 456 A (LUB JOHAN ET AL) 11 August 1998 (1998-08-11) cited in the application abstract claims 1,2	19,20
A	US 5 691 789 A (FARIS SADEG MUSTAFA ET AL) 25 November 1997 (1997-11-25) figures 3,5	4,5,19, 20
	-/-	

☒ Further documents are listed in the continuation of box C.☒ Patent family members are listed in annex.

* Special categories of cited documents:

"A" document defining the general state of the art which is not considered to be of particular relevance

"E" earlier document but published on or after the international filing date

"L" document which may throw doubts on priority claim(s) or which is cited to establish the publication date of another citation or other special reason (as specified)

"O" document referring to an oral disclosure, use, exhibition or other means

"P" document published prior to the international filing date but later than the priority date claimed

"T" later document published after the international filing date or priority date and not in conflict with the application but cited to understand the principle or theory underlying the invention

"X" document of particular relevance; the claimed invention cannot be considered novel or cannot be considered to involve an inventive step when the document is taken alone

"Y" document of particular relevance; the claimed invention cannot be considered to involve an inventive step when the document is combined with one or more other such documents, such combination being obvious to a person skilled in the art.

"G" document member of the same patent family

Date of the actual completion of the international search

30 March 2000

Date of mailing of the international search report

06/04/2000

Name and mailing address of the ISA

European Patent Office, P.B. 5818 Patentlaan 2
NL - 2280 HV Rijswijk
Tel. (+31-70) 340-2040, Tx. 31 651 epo nl,
Fax: (+31-70) 340-3018

Authorized officer

Gill, R

INTERNATIONAL SEARCH REPORT

Int. Application No

PCT/EP 99/09711

C.(Continuation) DOCUMENTS CONSIDERED TO BE RELEVANT

Category *	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
E	<p>EP 0 982 605 A (MERCK PATENT GMBH) 1 March 2000 (2000-03-01) abstract page 4, line 15 - line 18 page 5, line 25 - line 31 page 6, line 10 - line 18 claims 2,10</p>	15-20

INTERNATIONAL SEARCH REPORT

Information on patent family members

International Application No

PCT/EP 99/09711

Patent document cited in search report	Publication date	Patent family member(s)	Publication date
US 5668614 A	16-09-1997	NONE	
US 5793456 A	11-08-1998	AT 178996 T CN 1095164 A DE 69417776 D DE 69417776 T EP 0606940 A JP 6281814 A US 5506704 A	15-04-1999 16-11-1994 20-05-1999 07-10-1999 20-07-1994 07-10-1994 09-04-1996
US 5691789 A	25-11-1997	AU 7601496 A CN 1217796 A EP 0873537 A JP 11512849 T WO 9716762 A US 6034753 A US 5940150 A	22-05-1997 26-05-1999 28-10-1998 02-11-1999 09-05-1997 07-03-2000 17-08-1999
EP 0982605 A	01-03-2000	NONE	